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(54) Alkylicarbamylmethylated aminotriazine crosslinking agents and curable compositions containing the same.

(57) Novel alkylicarbamylmethyl aminotriazines self-cure and also function as crosslinkers for compounds containing active hydrogen groups. When the active hydrogen-containing compounds are hydroxylated polymer, coatings are provided with exceptional resistance to detergent and salt-spray exposure and improved abrasion resistance. The novel aminotriazines also can be combined with fillers as binders which when cured provide shaped articles of manufacture, such as insulation and foundry core molds.

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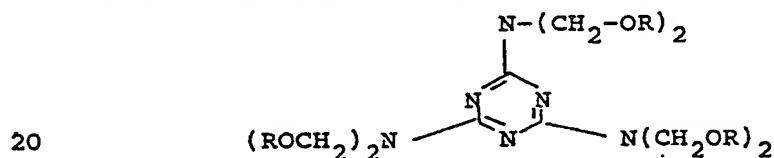
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**ALKYLCARBAMYL METHYLATED AMINOTRIAZINE
CROSSLINKING AGENTS AND CURABLE
COMPOSITIONS CONTAINING THE SAME**

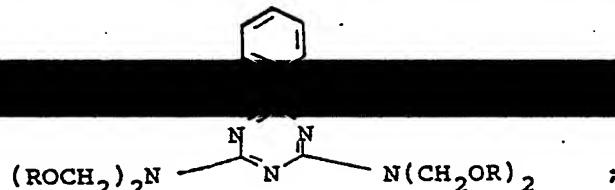
The present invention relates to curing agents, to curable compositions and to methods of making and using them. More particularly, the present invention relates to novel curing agents comprising alkylcarbamylmethylaminotriazines and to curable compositions comprising an active hydrogen-containing material, a novel alkylcarbamylmethylaminotriazine and a cure catalyst. Coatings cured from the compositions have exceptional resistance to detergent and salt spray exposure, making them well adapted for use in powder coatings, coil coatings and can coatings. The new compositions can be used with filler to provide shaped articles of manufacture with superior properties.

BACKGROUND OF THE INVENTION

Curable compositions containing aminotriazine compounds are known in the art. As is shown in Koral et al., U.S. 3,661,819, for example, a preferred family of aminotriazine curing agents comprises (i) a triamino-triazine compound of the formula:



which will be depicted hereinafter as $\text{C}_3\text{N}_6(\text{CH}_2\text{OR})_6$; or (ii) a benzoguanamine compound of the formula:



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which will be depicted hereinafter as $C_3H_5(C_6H_5)(CH_2OR)_4$ wherein R is hydrogen or alkyl of from 1 to 12 carbon atoms. It is also known to use oligomers of such compounds, which are low molecular weight condensation products containing
5 for example two, three or four triazine rings, joined by $-CH_2OCH_2-$ linkages, as well as mixtures of any of the foregoing. These are used to cure active hydrogen-containing materials, especially polymers which contain carboxyl groups, alcoholic hydroxy groups, amide groups and groups
10 convertible to such groups, such as methylol groups. When such curable compositions are applied to substrates as coatings and then cured, excellent hardness, impact resistance, light stability and solvent resistance is imparted to the articles. The compositions can also be formulated with
15 fillers and/or reinforcements such as particulate and fibrous mineral and organic materials, such as cellulose, wood, glass, graphite, textiles, silica, asbestos, wollastonite, and the like to produce insulation, foundry molds and the like which have superior properties and show a
20 reduced tendency to emit formaldehyde during use.

As is described in German Patent OL 2,005,693 (1971) (Chemical Abstracts 76:P 34864 a (1972)), when triaminotriazines of the general formula (i) above are reacted with arylurethanes, such as phenyl urethane, there are
25 produced reaction products of the typical formula $C_3N_6(CH_2-NH-COOC_6H_5)_6$, and when these are reacted with polymers containing hydroxyl groups such as acrylics and polyesters, crosslinking occurs with the development of colorless, very hard films, which remain colorless even when the baking time
30 is increased tenfold, to five hours at 100°C. However, subsequent experiments have shown that such coatings, like those crosslinked with the triazines of formulae (i) and (ii) above, are somewhat deficient in detergent resistance, salt spray resistance and adhesion. They also are produced
35 with the liberation of phenol, which causes health and

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disposal problems, and is economically wasteful.

It has now been discovered that if aminotriazines of general formulae (i) and (ii) are reacted with alkylurethanes (which are well known to be less reactive than 5 the aryl carbamates used in the above-mentioned German Patent), derivatives are formed which are also reactive to crosslink active hydrogen-containing polymers, but the new coatings which are formed have much improved properties (detergent, salt spray, adhesion, color retention) over 10 those of the prior art, particularly the aryl-substituted derivatives of OL 2,005,693.

Although it is known, e.g., from Amin et al., Indian J. Chem., 14B, 139-140 (1976), to prepare both aryl and alkyl carbamylmethylated melamines, by the 15 reaction of trimethylolmelamine with n-hexyl carbamate, only the mono-substituted product was produced, and this would not be capable of acting as a crosslinker to introduce two or more urethane groups. Such groups are now believed to be essential to secure all of the 20 advantages of the present invention.

Thus the present invention differs from the state of the art by providing aminotriazine derivatives containing at least two alkylcarbamylmethyl groups and then using them as crosslinkers for active hydrogen-containing materials to 25 provide coatings with exceptional resistance, for example, to detergent and salt-spray exposure, and improved light stability.

SUMMARY OF THE INVENTION

According to the present invention there are 30 provided triazine compounds selected from:

(i) a triaminotriazine compound of the formula

35 $C_3N_5(C_6H_5)(CH_2OR)^{4-y}(CH_2NHCOOR^1)_y$;
 (iii) an oligomer of (i) or of (ii) ; or

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(iv) a mixture of at least two of any of (i), (ii) and (iii), wherein the R groups are, independently, hydrogen or alkyl of from 1 to 12 carbon atoms, the R¹ groups are, independently, alkyl of from 1 to 20 carbon atoms, x is in the range of from about 2 to about 6, and y is in the range of from about 2 to about 4;

In preferred embodiments of the invention, x is in the range of from about 2.8 to about 6 and y is in the range of from about 2.2 to about 4. With respect to compound (i) R is lower alkyl, preferably C₁-C₈ and R¹ is methyl, ethyl, n-propyl, i-propyl, butyl, n-octyl, 2-ethylhexyl, n-octadecyl, or a mixture of any of the foregoing. Also preferred are oligomers of (iii)(i) in which R is methyl and R¹ is methyl, ethyl, n-propyl, i-propyl, butyl or a mixture of any of the foregoing as well as benzoguanamines (ii) wherein R and R¹ are the same as defined above with respect to compound (i).

Also contemplated by the present invention are curable compositions comprising

- 20 (a) an active hydrogen-containing material;
- (b) a triazine compound selected from
 - (i) a triaminotriazine compound of the formula C₃N₆(CH₂OR)_{6-x}(CH₂NHCOOR¹)_x;
 - (ii) a benzoguanamine compound of the formula C₃N₅(C₆H₅)(CH₂OR)_{4-y}(CH₂NHCOOR¹)_y;
 - (iii) an oligomer of (i) or of (ii); or
 - (iv) a mixture of at least two of any of (i), (ii) and (iii), wherein the R groups are, independently, hydrogen or alkyl of from 1 to 12 carbon atoms,
- 30 the R¹ groups are, independently, alkyl of from 1 to 20 carbon atoms, x is in the range of from about 2 to about 6, and y is in the range of from about 2 to about 4; and
- (c) a cure catalyst.

In preferred features of this aspect of the invention, the active-hydrogen containing material (a) is a

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polymeric material containing at least two reactive carbonyl, alcoholic hydroxy, amide or amine groups, or a mixture of such groups, or a group convertible to such groups, preferably a hydroxy-functional acrylic resin or a low molecular weight polyester polyol. Preferably the triazine will be as set forth specifically above, and the cure catalyst will be a metal salt or metal complex comprising tin, especially preferably tetrabutyldiacetoxy stannoxane.

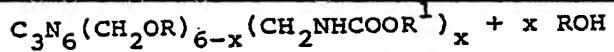
10 Alternatively, the alkylcarbamylmethyl triazines can be used as a self-crosslinkable material in providing protective and/or decorative coatings.

15 Also provided by the invention are articles of manufacture comprising substrates protectively coated with a cured composition as defined above and articles of manufacture comprising a cured composition as defined above and a filler, e.g., glass, e.g., glass powder, glass beads, glass fibers or foundry sand.

DETAILED DESCRIPTION OF THE INVENTION

20 As starting materials to produce the alkylcarbamylmethylated triazines of this invention, there can be used the hydroxymethyl or alkoxyethyl melamines and/or benzoguanamines and oligomers thereof known in the art. Many of the starting materials are commercially available, 25 and can be made by well known procedures. In accordance with the present invention, the starting materials are reacted with alkyl carbamates, such as methyl carbamate and propyl carbamate, which also are well known in this art, in the presence of an acid catalyst.

30 An idealized reaction equation for the preparation of the new compounds from an alkoxyethylmelamine or a hydroxymethylmelamine is as follows:



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wherein R, R¹ and x are as defined above.

The mole ratio of alkyl carbamate is selected to provide the desired degree of substitution. By way of illustration, from 2 to 6 moles can be used. Reaction is

- 5 typically carried out by heating in the melt or in solution, e.g., in benzene, toluene, xylene, chlorobenzene, dichlorobenzene, e.g., in the presence of catalytic amounts of acid, e.g., para-toluenesulfonic acid, nitric acid, sulfuric acid, and the like, at temperatures between 80°
- 10 and 150°C., preferably 90-120°C. Measurement of the quantity of alcohol (ROH) evolved gives an indication of reaction completion. With 6 moles of alkyl carbamate, reaction is usually not 100% complete, unless forced, but a high degree of substitution, x = 5-6, is obtained.
- 15 Analysis by gel permeation chromatography shows that treatment of hexamethoxymethylolmelamine with substantially less than 6 moles of alkyl carbamate gives a product distribution similar to the starting material with degrees of substitution ranging up to 6. Of course,
- 20 only those compounds wherein at least two carbamylmethyl groups are present are crosslinkers according to this invention, even though residual alkoxyethyl groups can provide crosslinking.

Instead of alkoxyethylmelamines, hydroxymethylmelamines, and the corresponding benzoguanamine analogs and oligomers can be used as starting materials. The products can be recovered by any convenient means after removal of byproduct water or alcohol is complete. Simply cooling to room temperature will leave the product as a residue, and the acid catalyst can be removed by neutralization.

The substituents defined by R and R¹ in the formulae above can vary widely in carbon content, and the groups can be straight chain, branched chain and alicyclic. A number of representative compounds will be exemplified in

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detail hereinafter.

The active hydrogen-containing materials have as the active hydrogen group a group selected from carboxyl, alcoholic hydroxyl, amido, primary amine, secondary amine 5 (including imine), thiol and the like. The active hydrogen-containing materials useful herein are typically film-forming compositions. Illustrative examples of active hydrogen-containing materials are shown in the above-mentioned Koral patent, the above-mentioned German OLS 10 2,055,693, and in Valko, U.S. 4,435,559. Typical polymers are acrylic polymers, polyesters, epoxy resins, and the like, providing that they contain active hydrogen groups.

Especially suitable are polyesters and polyacrylates containing pendant hydroxyl groups as reaction sites. 15 The former are obtained in a known manner by the reaction of polycarboxylic acids with excess quantities of polyhydric alcohols; the latter are obtained by the copolymerization of acrylic or methacrylic acid derivatives with hydroxyl-group-containing derivatives of these acids, such 20 as, for example, the hydroxyalkyl esters, optionally with the simultaneous use of additional vinyl compounds, such as, for example, styrene. Hydroxyl-group-containing polyurethanes can be obtained in known manner by the reaction of polyisocyanates with excess quantities of compounds 25 containing at least two hydroxy groups. Suitable commercially available hydroxy-group-containing polyesters are CYPLEX® 1473 and CYPLEX® 1531 from American Cyanamid Company and Cargil Polyester 5776. Suitable hydroxy-functional acrylic resins are available commercially from S.C. Johnson 30 & Son, Inc. under the trademark JONCRYL®-500. Also suitable for use are a hydroxy-terminated polycaprolactone, as well

35 mentioned German OLS 2,055,693 and the polyester of phthalic acid, adipic acid, ethanediol, and trimethylolpropane, with

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a hydroxy number of 130 and an acid number of 1.5 of Example 6 of the said OLS publication.

As set forth herein, the curable composition includes a cure catalyst. Typically, the cure catalyst is a metal salt and/or complex of a metal such as lead, zinc, iron, tin and manganese, preferably tin. Suitable salts of these metals are, for example acetates, octoates, laurates and naphthanates. Suitable complexes, for example, are tetrabutyldiacetoxy stannoxane, dibutyltin dilaurate, dimethyltin dilaurate or an acetyl acetonate. The cure catalyst is used in amounts effective to accelerate cure at the temperatures employed, e.g., 120-220°C. For example, the catalyst is used in amounts from about 0.1 to about 2.0 preferably 0.2 to 1% metal by weight (solids) based on the weight of the curable compositions.

It should also be understood that residual ether functional groups can cure with catalysts usually used with amino resins, such as acid catalysts, e.g., nitric acid, sulfuric acid, p-toluenesulfonic acid and the like. This may be advantageous where lower cure temperatures are useful, e.g., when binding fillers or reinforcements, e.g., textiles, cellulose, wood flour, etc. Also useful as heterogenous acidic catalysts are ion exchange resins in the acid form.

In the practice of the invention, the curable compositions can be adapted for use in solvent-based or water based coating compositions. Coating compositions comprising aqueous dispersions are particularly suited to application by electrodeposition. Generally the compositions will contain about 1 to 75 percent by weight of resin and crosslinker combined, and the weight ratio of crosslinker to resin will range from about 5 to about 40 parts to correspondingly from 60 to 95 parts of said resin.

In many instances a pigment composition and various conventional additives such as antioxidants, surface active

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agents, coupling agents, flow control additives, and the like, can be included. The pigment composition may be of any conventional type, such as, one or more pigments such as iron oxides, lead oxides, strontium chromate, carbon black,

5 titanium dioxide, talc, barium sulfate, cadmium yellow, cadmium red, chromic yellow, or the like.

After deposition on a substrate, such as a steel panel, the coating composition is devolatilized and cured at elevated temperatures by any convenient method such as

10 in baking ovens or with banks of infrared heat lamps. Curing can be obtained at temperatures in the range of from 120°C. to about 300°C., preferably from 150°C. to about 200°C. for from about 30 minutes at the lower temperatures to about 1 minute at the higher temperatures.

15 Conventional methods can be used to combine the novel aminotriazines herein with fillers and/or reinforcements and to shape them into useful articles by means well known to accomplish these functions with curable aminotriazine resins. Mixing with glass fillers for example and

20 heating provides insulation shapes for pipes, and the like, after curing, and mixing with foundry sand and curing provides core molds for metal casting. These have superior strength compared to the state of the art and appear to be highly advantageous in not evolving formaldehyde during use.

25 DESCRIPTION OF THE PREFERRED EMBODIMENTS

The following examples illustrate the compounds and compositions of the present invention. They are not to be construed as limiting the claims in any manner. All parts are by weight.

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EXAMPLE 1

Reaction Product of 6 Moles of N-
Propyl Carbamate With 1 Mole of
Hexamethoxymethylmelamine (6-PC)

5 Hexamethoxymethylmelamine (47.2g, 0.121 mole, American Cyanamid Co. CYMEL® 300), n-propylcarbamate (75.0 g, 0.728 mole), and para-toluenesulfonic acid (0.33 g) are stirred at 95°C. in a flask equipped with a vacuum distillation head. During 50 minutes, the
 10 pressure is lowered in stages to 50 mm Hg and 21.1 g of methanol (0.659 mole, 91% of theoretical) is collected in the distillate receiver. The product in the reaction flask is cooled to near room temperature, where it is a clear, colorless, very viscous liquid. Methylene
 15 chloride (100 ml) is added and stirring for one-half hour dissolves the product. The acid catalyst is removed by washing with sodium carbonate solution, followed by drying over potassium carbonate. Rotary vacuum evaporation gives 98.6 g of clear, colorless,
 20 nearly solid product. Nuclear magnetic resonance (NMR) analysis shows that the product has at least five (on average) of the methoxy groups replaced by n-propyl carbamate groups:

25 $C_3N_6(CH_2OCH_3)_{0-1}(CH_2NHCOOCH_2CH_2CH_3)_{5-6}$
 Gel permeation chromatography shows one large peak for the monomeric compound (> 80%) and two smaller peaks corresponding to dimeric (~10%) and trimeric (~3%) oligomers.

COMPARATIVE EXAMPLE 1A

30 Reaction Product of 6 Moles of Phenyl Carbamate With 1 Mole of
Hexamethoxymethylmelamine (6-PhC)

For comparison purposes, the procedure of Example 1 of German OLS 2,005,693 is repeated: Hexamethoxymethyl-melamine (300 g, 1 mole) and 822 grams of phenyl-

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urethane (6 moles) are dissolved in two liters of chlorobenzene and, after the addition of 4 grams of p-toluene-sulfonic acid, are heated with a vertical condenser, with stirring and passage of CO₂, until the boiling point of 5 chlorobenzene is reached. The methanol cleavage starts at approximately 90 to 100 C. A mixture of methanol with a small amount of chlorobenzene comes over first. With increasing temperature, the boiling point of chlorobenzene (130°C) is reached.

10 The mixture is then evaporated under vacuum, with stirring, to produce a colorless resin with a softening point of 85 to 120°C, which is soluble in all proportions in ethyl acetate. The yield is high. The product is of the formula C₃N₆(CH₂NHCOOC₆H₅)₆.

15 EXAMPLE 2

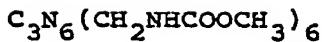
Reaction Product of 6 Moles of
Methyl Carbamate with One Mole of
Hexamethoxymethylmelamine (6-MC)

Hexamethoxymethylmelamine (19.5 g, 0.05 mole), 20 excess methyl carbamate (37.6 g, 0.50 mole), and para-toluene-sulfonic acid (0.86 g, 0.005 mole) are stirred at 97°C. in a flask equipped with a distillation head as in Example 1. The reaction mixture changes from a clear, colorless liquid to a white solid and a few ml. 25 of distillate is formed. The reaction mixture is then allowed to cool to room temperature.

A portion, 20.0 g, of the solid product is powdered and vigorously stirred with 100 ml. of water at room temperature for 1½ hours. Filtration and drying 30 gives 12.3 g of white solid; m.p. = 179-188°C. Infrared spectroscopy shows that at least 90-95% of the methoxy groups have been replaced by carbamate groups.

theoretical yield of hexamethylcarbamylmethylated melamine,
35 a compound of the formula

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EXAMPLE 3

Reaction Product of 2 Moles of n-Octadecyl Carbamate With 1 Mole of Hexamethoxymethylmelamine

5 Hexamethoxymethylmelamine (25.0 g, 0.0641 mole), n-octadecyl carbamate (40.1 g, 0.128 mole), and 0.31 g (0.0018 mole) of para-toluenesulfonic acid are stirred at 100°C in a flask equipped with a vacuum distillation head.

10 During 30 minutes, the pressure is lowered in stages to 50 mm Hg and distillate is collected in the receiver.

At room temperature the product is opaque, white and has the consistency of mayonnaise. Gas chromatographic analysis of a sample of the product dissolved in 15 methyl isobutyl ketone shows practically no unconverted octadecyl carbamate. The formula is, approximately:



EXAMPLE 4

20 Reaction Product of 6 Moles of Isopropyl Carbamate With 1 Mole of Hexamethoxymethylmelamine

Hexamethoxymethylmelamine (18.9 g, 0.0485 mole), isopropyl carbamate (30.0 g, 0.291 mole), and para-toluene-sulfonic acid (0.13 g., 0.008 mole) and stirred at 95°C 25 in a flask equipped with a vacuum distillation head. During 45 minutes, the pressure is lowered in stages to 50 mm Hg, and 8.46 g of methanol (0.264 mole, 91% of theoretical) is collected in the distillate receiver. The product is dissolved in 75 ml of methylene chloride and the solution 30 is washed with two portions of aqueous 5% sodium carbonate to remove the acid catalyst. The solution is dried over anhydrous potassium carbonate and rotary vacuum evaporated to give 36.8 g of colorless solid (93% yield). The solid is pulverized to white powder; m.p. 101 to 130°C 35 (clear, colorless melt). The formula is, approximately:

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EXAMPLE 5

Reaction Product of 2 Moles of n-Octyl Carbamate With 1 Mole of Hexamethoxymethylmelamine

5 Hexamethoxymethylmelamine (22.6 g, 0.0579 mole), n-octyl carbamate (20.0 g, 0.116 mole), and para-toluene-sulfonic acid (0.19 g, 0.0011 mole) are stirred at 75°C in a flask equipped with a vacuum distillation head. During
 10 50 minutes, the pressure is lowered in stages to 50 mm Hg, and 3.58 g of methanol (0.112 mole, 96% of theoretical) is collected in the distillate receiver. The product is dissolved in 150 ml of methylene chloride and the solution is washed with two portions of aqueous 5% sodium carbonate
 15 to remove the acid catalyst. The solution is dried over anhydrous potassium carbonate and rotary vacuum evaporated to give 37.2 g of almost-clear, colorless, viscous liquid (96% yield of theoretical product). The infrared spectrum is consistent with the expected structure. Gel permeation
 20 chromatography shows peaks attributed to mono-, di-, tri-, and higher-substituted products; little oligomeric material is evident. The formula is, approximately:

EXAMPLE 6

25 Reaction Product of 6 Moles of n-Propyl Carbamate With 1 Mole of Hexamethylolmelamine

The general procedure of Example 1 is repeated, substituting the hydroxymethyltriazine: Hexamethylol-
 30 melamine (10.0 g, 0.0327 mole), n-propyl carbamate (20.2 g, 0.196 mole), and para-toluenesulfonic acid (0.60 g, 0.0035 mole) are stirred at 95°C in a flask equipped with

35 distillate (mostly water, 92% of theoretical) is collected

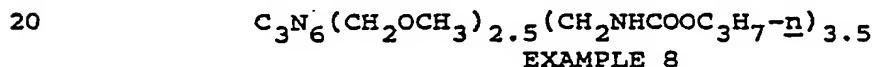
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in the distillate receiver. At room temperature, the product in the reaction vessel is a gray-white solid. The infrared spectrum shows little or no residual hydroxyl functionality and is similar to the spectrum of authentic 5 hexa-n-propylcarbamylmethylated melamine, made by Example 1. The general formula is, approximately:



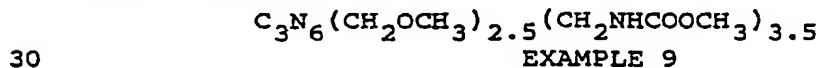
10 Reaction Product of 3.5 Moles of
n-Propyl Carbamate with 1 Mole
of Hexamethoxymethylmelamine

The general procedure of the preceding Examples is used to react hexamethoxymethylmelamine (167.8 g, 0.430 moles) with 155.1 g, 1.566 moles of n-propyl carbamate. The acid catalyst in this instance is 0.7 g of concentrated nitric acid. The product weighs 254.2 g and contains only 0.1% residual carbamate. It melts at 85-15 95°C. A 65% solids solution in xylene remains clear and colorless for 8 weeks. It has the formula, approximately:



Reaction Product of 3.5 Moles of
Methyl Carbamate With 1 Mole of
Hexamethoxymethylmelamine

25 The procedure of Example 7 is repeated substituting 1.589 moles of methyl carbamate. The product weighs 241.7 g and melts at 95-103°C. It has the formula, approximately:



Reaction Product of 3.5 Moles of n-Propyl Carbamate With 1 Mole of
Hexamethoxymethylmelamine Oligomer

35 The procedure of Example 7 is repeated, substituting an oligomeric triazine (American Cyanamid Co.

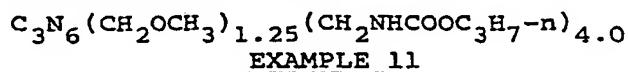
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CYTEL® 303), $C_3N_6(CH_2OCH_3)_{5.25}$. The catalyst is removed by extracting a xylene solution of the product mixture with sodium carbonate solution. A 75% solution in xylenes remains clear and colorless for more than 6 weeks. The product melts at 74-80°C. The formula is, approximately:



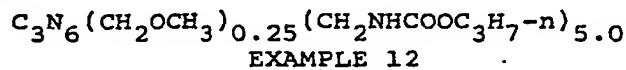
10 Reaction Product of 4 Moles of n-Propyl Carbamate With 1 Mole of Hexamethoxymethylmelamine Oligomer

The procedure of Example 9 is repeated with the higher mole ratio of n-propyl carbamate. The product, 412.2 g, melts at 80-90°C. At 75% solids in xylenes and 15 methyl isobutyl ketone, the product in solution remains clear and colorless for more than 6 weeks. It has the following approximate formula:



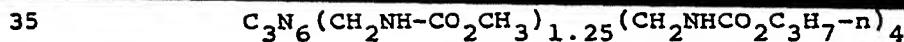
20 Reaction Product of 5 Moles of n-Propyl Carbamate With 1 Mole of Hexamethoxymethylmelamine Oligomer

The procedure of Example 9 is repeated with a higher mole ratio of n-propyl carbamate. The product, 25 32.4 g, melts at 80-92°C. It has the following approximate formula:



30 Reaction Product of 4 Moles of n-Propyl Carbamate and 2 Moles of Methyl Carbamate With 1 Mole of Hexamethoxymethylmelamine

The general procedure of Example 9 is used at 35 °C. The product, 95.8 g, melts at 85-95°C. The formula

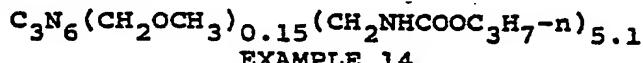


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EXAMPLE 13

Reaction Product of 5.25 Moles of
n-Propyl Carbamate With 1 Mole of
Hexamethoxymethylmelamine Oligomer

5 The general procedure of Example 9 is used. The product weighs 113.5 g and melts at 90-100°C. The formula is, approximately:

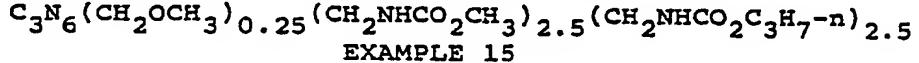


EXAMPLE 14

10 Reaction Product of 2.5 Mole of n-Propyl Carbamate and 2.5 Mole of Methyl Carbamate With 1.0 Mole of
Hexamethoxymethylmelamine Oligomer

The general procedure of Example 9 is used at 110°C. The product weighs 174.5 g and melts at 85-95°C.

15 The general formula is, approximately:



EXAMPLE 15

20 Reaction Product of 3.5 Mole of n-Propyl Carbamate with 1 Mole of (Trimethoxy-methyl-Tributoxymethyl) Melamine
(Trimethoxymethyl-tributoxymethyl)melamine
(CYMEL® 1133, American Cyanamid Co., 320.3 g, 0.689 mole), n-propyl carbamate (248.3 g, 2.411 mole) and 1.12 g of concentrated nitric acid are stirred and heated in

25 a 100°C. oil bath under a steady stream of nitrogen. Distillate is collected in a dry-ice/isopropanol cooled trap. After 60 minutes, during which vacuum of up to 50 mm Hg is applied, the reaction is stopped. The hot, crude reaction product is dissolved to 30% solids in

30 mixed xylenes (1059 g of xylenes are added) and extracted once with 300 ml of 5% aqueous sodium carbonate solution. The organic layer is then extracted several times with hot deionized water to neutralize the acid and reduce the amount of residual n-propyl carbamate.

35 The organic layer is dried over potassium carbonate

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(anhydrous) until clear and then stripped under vacuum of a 65% solids content. The product weighs 698.5 grams.

EXAMPLE 16

5 Reaction Product of 3.5 Mole of Methyl Carbamate with 1 Mole of (Trimethoxy-methyl-Tributoxymethyl)Melamine

The procedure of Example 15 is repeated, substituting 376.28 g, 0.809 mole of trimethoxymethyl-
10 tributoxymethylmelamine, 212.42 g, 2.832 mole of methyl carbamate and 1.32 g of conc. nitric acid. The product is recovered by the procedure of Example 9.

In the following examples, the alkylcarbamylmethylated triazines of this invention are formulated
15 into curable compositions and evaluated as coatings. For comparison purposes, the phenylcarbamylmethylated triazine of German OLS 2,005,693 (Example 1A herein) is also evaluated.

The general method of preparation is as follows:
20 Thermosetting coatings containing polyols with alkyl and phenylcarbamylmethylated melamines on steel are prepared by mixing high solids solutions of alkylcarbamyl and arylcarbamylmethyl melamines (65% solids in methyl isobutyl ketone or xylene) with a solution of a hydroxyl
25 functional acrylic resin (specifically, JONCRYL®-500, S.C. Johnson & Son, Inc.; 85% solids in methyl amyl ketone), a solution of a tin catalyst (tetrabutyldiacetoxy stannoxyane, TBDSA, 10% solids in methyl isobutyl ketone), and a flow control additive (FC-431, 10% solids in ethyl acetate, 3M
30 Co.). A second series of coatings compositions is prepared as above but with low molecular polyester polyol as backbone resin (specifically, CYPLEX® 1473, 60% solids in

lated with and without EPON®-1001, an epoxy resin (10

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parts per hundred-phr, as an 85% solids solution in methyl isobutyl ketone - Shell Co.) to assess resistance properties in particular.

The coatings are applied using #40 or #46 WIRE-
 5 CATORS® by drawdown of resin formulations on 4" x 12"
 BONDERITE®-100 treated steel panels. The panels, after
 drawdown, are held for 10 minutes at room temperature and
 then cured on racks in a temperature controlled oven, at
 specified temperatures. The coatings prepared are about
 10 1.2 ± 0.2 mils thick and initially tested for solvent
 resistance by rubbing with a cloth saturated with methyl
 ethyl ketone (MEK rubs) in accordance with standard test-
 ing methods.

Table 1 illustrates a typical charge composition
 15 for preparing a coated panel.

TABLE 1

<u>Material</u>	<u>% Solids</u>	<u>PHR*</u>	<u>Charge (Grams solution)</u>
CYPLEX® 1475-5	65	70	32.3
<u>Propyl Carbamate</u>			
20 Melamine resin	60	20	10.0
EPONE®-1001	75	10	4.0
TBDAS cat.	10	1	3.0
FC-431	10	0.13	0.4

* PHR = parts per hundred resin; final solution 57.8% solids.

25 Materials 1--5 are stirred until homogeneous, then
 filtered through a 10 micron felt filter to remove small
 particles and deaerated.

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The properties of the coatings evaluated include:

<u>Property</u>	<u>Method</u>
Forward and Reverse Impact	ASTM D-3281-73
Color	Visual
5 20° Gloss	Measured on Glossgard II 20°/60° Gossmeter - Neotec Instr. Div., Pacific Scientific
Detergent Resistance at 72°C.	ASTM D-2248-73; reapproved
10	1982; Evaluation of Degree of Blistering of Paints D-714
Blister Classification	Example: F-8 means few small blisters; D-4 means dense large blisters. The smaller the number following the letter, the larger the blister on a scale of 1-10, with 10 meaning <u>no</u> blistering.
15	
20	

EXAMPLES 17-22

The crosslinker of Example 1 herein, the reaction product of 6 moles of n-propyl carbamate and 1 mole of hexamethoxymethyl melamine (6-PC), is used with a hydroxy-functional polyacrylate and tetrabutyldiacetoxystannoxane as cure catalyst. For comparison purposes, formulations are made substituting the reaction product of 6 moles of phenyl carbamate with hexamethoxymethyl melamine (6-PhC) of Comparative Example 1A. The formulations used and the properties of the cured films are set forth in Table 2:

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TABLE 2: Carbamylmethylated Melamine Crosslinked Acrylic Resin

<u>Example</u>	<u>17</u>	<u>18</u>	<u>19</u>	<u>20</u>	<u>21</u>	<u>22</u>
<u>Composition (parts by weight)</u>						
5 Polyacrylate ^a	80	70	60	50	80	60
6-PC ^b	20	20	40	40	20	40
6-PhC ^c	--	--	--	--	--	--
Epoxy Resin ^d	--	10	--	10	--	--
TBDAS ^e	1	1	1	1	1	1
10 Cure Temp., °C.	180	180	180	180	160	160
Cure Time, min.	20	20	20	20	30	30
<u>Properties</u>						
MEK wipes	200+	200+	200+	200+	200+	200+
Thickness, mils.	1.2	1.1	1.0	1.0	1.2	1.1
15 Forward Impact,	30	10	10	20	20	--
Color	Clr	Clr	Clr	Clr	Clr	Clr
20° Gloss	97	93	95	93	97	96
Knoop Hardness	13.5	13	16	15.8	10.8	16.2
<u>Detergent Immersion</u>						
20 72 hrs	M8	10	F9	10	M8	D9
120 hrs	D8	10	F8	10	M6	D8
192 hrs	D8	D9	F9	10	D7	M9
240 hrs	--	--	--	10	--	--
288 hrs	--	--	--	10	--	--
25 408 hrs	--	--	--	10	--	--
456 hrs	--	--	--	10	--	--
744 hrs	--	--	--	10	--	--
912 hrs	--	--	--	10	--	--
1100 hrs	--	--	--	10	--	--

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TABLE 2: CONTINUED

	<u>Example</u>	<u>17</u>	<u>18</u>	<u>19</u>	<u>20</u>	<u>21</u>	<u>22</u>
<u>Composition (parts by weight)</u>							
	Polyacrylate ^a	80	70	60	50	80	60
5	6-PC ^b	--	--	--	--	--	--
	6-PhC ^c	20	20	40	40	20	40
	Epoxy Resin ^d	--	10	--	10	--	--
	TBDAS ^e	1	1	1	1	1	1
	Cure Temp., °C.	180	180	180	180	160	160
10	Cure Time, min.	20	20	20	20	30	30
<u>Properties</u>							
	MEK wipes	--	200 ^m	200+	200+	200 ^m	--
	Thickness, mils.	1.1	0.94	0.88	0.83	1.2	1.0
	Forward Impact,	--	20	20	20	10	--
15	Color	Clr	Yel	Sl.Brn	Yel	Sl.Yel	Clr
	20° Gloss	94	97	95	91	92	86
	Knoop Hardness	15.3	13	18	17	12.4	19.4
<u>Detergent Immersion</u>							
	72 hrs	M6	10	F6	10	VD6	M8
20	120 hrs	D2	M8	F8	10	VD4	M8
	192 hrs	--	D8	F8	10	--	M8
	240 hrs	--	--	--	10	--	--
	288 hrs	--	--	--	10	--	--
	408 hrs	--	--	--	10	--	--
25	456 hrs	--	--	--	F6	--	--
	744 hrs	--	--	--	F6	--	--
	912 hrs	--	--	--	F6	--	--
	1100 hrs	--	--	--	F6	--	--

30 a JONCRYL® S.C. Johnson & Son, Inc., 85% solids in methyl-
amylketone;

b n-propylcarbamylmethyl melamine (Example 1);

c phenylcarbamylmethyl melamine (Comp. Ex. 1A);

35 m = marred; and s = softened

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EXAMPLES 23-30

The crosslinkers of Examples 7 and 8 herein (3.5 PC and 3.5 MC) are used to crosslink the hydroxy functional acrylic resin and evaluated in coatings. The formulations used and the properties obtained are set forth in Table 3:

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TABLE 3: Carbamylmethylated Melamine-Crosslinked Acrylic Resins

<u>Example</u>	<u>23</u>	<u>24</u>	<u>25</u>	<u>26</u>	<u>27</u>	<u>28</u>	<u>29</u>	<u>30</u>
<u>Composition (parts by weight)</u>								
	Polyacrylate ^a	80	70	60	50	80	70	60
5	3.5-PC ^b	20	20	40	40	--	--	--
	3.5-MC ^c	--	--	--	--	20	20	40
	Epoxy Resin ^d	--	10	--	10	--	10	--
	TBDAS ^e	1	1	1	1	1	1	1
	Cure Temp., °C.	180	180	180	180	160	160	180
10	Cure Time, min.	20	20	20	20	20	20	20
<u>Properties</u>								
	MEK wipes	200+	200+	200+	200+	200+	200 ^m	200+
	Thickness, mils.	1.0	1.1	1.2	1.1	1.0	1.0	0.95
	Forward Impact,	30	5	20	18.5	30	15	15
15	Color	Clr	Clr	Clr	Clr	Clr	Clr	Clr
	20° Gloss	90	95	95	91	97.5	99.5	101
	Knoop Hardness	11.8	10.4	15.4	18.5	13	15	16.4
<u>Detergent Immersion</u>								
	48 hrs	M8	--	--	--	--	--	--
20	72 hrs	--	10	M8	10	M9	10	10
	120 hrs	D8	D8	M4	10	D6	10	10
	192 hrs	--	D8	D8	10	D8	10	10
	240 hrs	--	--	--	10	--	M8	F8
	288 hrs	--	--	--	10	--	--	F8
25	408 hrs	--	--	--	10	--	--	M6
	456 hrs	--	--	--	10	--	--	M4
	744 hrs	--	--	--	10	--	--	M4
	912 hrs	--	--	--	10	--	--	M4
	1100 hrs	--	--	--	10	--	--	10
30	1400 hrs	--	--	--	10	--	--	F9

a See footnote Table 2

b 3.5 propyl carbamylmethyimelamine (Ex. 7)

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EXAMPLES 31-34

The crosslinker of Example 1 herein, the reaction product of 6 moles of n-propyl carbamate and 1 mole of hexamethoxymethylolmelamine (6-PC) is used with a low 5 molecular weight hydroxyfunctional polyester and tetra-butyldiacetoxy stannoxane as cure catalyst. For comparison purposes, formulations are made substituting the reaction product of 6 moles of phenylcarbamate with hexamethoxy-methylmelamine (6-PhC) of Comparative Example 1A. The 10 formulations used and the properties of the cured films are set forth in Table 4:

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TABLE 4: Carbamylmethylated Melamine-Crosslinked Polyester Resin

	<u>Example</u>	<u>31</u>	<u>32</u>	<u>33</u>	<u>34</u>	<u>31A</u>
<u>Composition (parts by weight)</u>						
5	Polyester ^a	80	70	60	50	80
	6-PC ^b	20	20	40	40	--
	6-PhC ^c	--	--	--	--	20
	Epoxy Resin ^d	--	10	--	10	--
	TBDAS ^e	1	1	1	1	1
10	Cure Temp., °C.	180	180	180	180	180
	Cure Time, min.	20	20	20	20	20
<u>Properties</u>						
	MEK wipes	200 ^s	200+	200	200+	200 ^s
	Thickness, mils.	.98	1.0	.90	1.0	.92
15	Reverse impact,	160	--	40	--	160
	Forward Impact,	160	--	70	--	160
	Color	Clr	Clr	Clr	Clr	Clr/Yel
	20° Gloss	99	102	104	102	92
	Knoop Hardness	15.4	18.8	19.5	18.8	14.6
20	Detergent Immersion					
	48 hrs	--	10	--	10	--
	72 hrs	10	--	10	--	10
	96 hrs	10	--	10	--	M8
	168 hrs	--	10	10	10	--
25	216 hrs	M6	10	F8	10	D4
	336 hrs	--	M8	--	10	--
	398 hrs	M2	--	M4	--	--
	504 hrs	--	D4-6	M2	10	--
	652 hrs	--	--	--	F8	--

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TABLE 4: CONTINUED

<u>Example</u>		<u>32A</u>	<u>33A</u>	<u>34A</u>	<u>34B</u>	<u>34C</u>
<u>Composition (parts by weight)</u>						
	Polyester ^a	70	60	50	80	60
5	6-PC ^b	--	--	--	--	--
	6-PhC ^c	20	40	40	40	40
	Epoxy Resin ^d	10	--	10	--	--
	TBDAS ^e	1	1	1	1	1
	Cure Temp., °C.	180	180	180	160	160
10	Cure Time, min.	20	20	20	30	30
<u>Properties</u>						
	MEK wipes	--	200	200	200	200 ^m
	Thickness, mils.	1.1	.85	.85	.95	.65
	Reverse impact,	--	140	30	160	160
15	Forward Impact,	--	150	70	160	160
	Color	Clr	Yel	S1Yel	Clr	Clr
	20° Gloss	106	107	105	98	101
	Knoop Hardness	19.1	21	22.5	16.2	21
<u>Detergent Immersion</u>						
20	48 hrs	--	--	--	--	--
	72 hrs	10	M8	10	10	F8
	96 hrs	--	D8	10	10	M6
	168 hrs	--	--	10	--	--
	216 hrs	10	D4	F8	F9	D8
25	336 hrs	--	--	--	--	--
	398 hrs	10	--	D4	D2	D2
	504 hrs	--	--	--	--	--
	652 hrs	D8	--	--	--	--

a American Cyanamid Co., CYPLEX® 1473-5

30 b-f See footnote above, Table 2.

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EXAMPLES 35-44

The crosslinkers of Examples 7 and 8, (3.5-PC and 3.5-MC) are used with the low molecular weight hydroxyfunctional polyester and tetrabutyldiacetoxy-
5 stannoxane as cure catalyst. The formulations used and the results obtained are set forth in Table 5:

**TABLE 5: Carbamylmethylated
Melamine-Crosslinked
Polyester Resins**

<u>Example</u>	<u>35</u>	<u>36</u>	<u>37</u>	<u>38</u>	<u>39</u>
5 Composition (parts by weight)					
Polyester ^a	80	70	60	50	80
3.5-PC ^b	20	20	40	40	--
3.5-MC ^c	--	--	--	--	20
Epoxy Resin ^d	--	10	--	10	--
10 TBDAS^e	1	1	1	1	.1
Cure Temp., °C.	180	180	180	180	180
Cure Time, min.	20	20	20	20	20
Properties					
MEK wipes	200 ^m	200+	200 ^s	200+	200+
15 Thickness, mils.	.92	.9	.77	.96	.88
Reverse impact,	160	--	120	--	150
Forward Impact,	160	100	120	70	160
Color	Clr	Clr	Clr	Sl.Yl.	Clr
20° Gloss	96	85	95	90	99
20 Knoop Hardness	14.2	17.4	18.8	21	14.4
Detergent Immersion					
48 hrs	10	10	--	--	--
72 hrs	--	--	10	10	10
96 hrs	--	--	10	10	M8
25 168 hrs	M6	10	--	--	--
216 hrs	D2	M8	M6	10	D4
336 hrs	--	D2	--	--	--
398 hrs	--	--	D2	F8	--
504 hrs	--	--	--	F6	--

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TABLE 5: CONTINUED

<u>Example</u>	<u>40</u>	<u>41</u>	<u>42</u>	<u>43</u>	<u>44</u>
<u>Composition (parts by weight)</u>					
Polyester ^a	70	60	50	80	60
5 3.5-PC ^b	--	--	--	--	--
3.5-MC ^c	20	40	40	20	40
Epoxy Resin ^d	10	--	10	--	--
TBDAS ^e	1	1	1	1	1
Cure Temp., °C.	180	180	180	160	160
10 Cure Time, min.	20	20	20	30	30
<u>Properties</u>					
MEK wipes	200+	200+	200+	--	--
Thickness, mils.	1.2	.85	1.1	1.1	1.0
Reverse impact,	50	80	10	--	--
15 Forward Impact,	50	80	40	--	--
Color	S1.Y1.	Clr	S1.Y1.	Clr	Clr
20° Gloss	101	99	102	101	102
Knoop Hardness	20	22.5	23.5	18	17.2
<u>Detergent Immersion</u>					
20 48 hrs	--	--	--	--	--
72 hrs	10	10	10	10	10
96 hrs	10	M8	10	10	10
168 hrs	--	--	--	--	--
216 hrs	10	D4	10	F8	10
25 336 hrs	--	--	--	--	10
398 hrs	10	--	10	M4	haze
504 hrs	--	--	10	--	VD8
652 hrs	--	--	VF8	VD2	--

a See footnote above, Table 4

b -do- ; Table 3

c -do- ; Table 3

d -do-

e -do-

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A review of the data in the foregoing tables indicates that improved properties are obtained with the alkylcarbamylmethylated melamines of this invention in comparison with the phenylcarbamylmethylated melamine of the prior art. Particularly outstanding with the acrylic and polyester coatings are the excellent detergent resistance properties of the new alkylcarbamylmethylated melamines. For example, both the methyl and propyl carbamylmethylated melamines give over 1100 hours of blister free coatings in a detergent bath when formulated with acrylic polyol and Epon-1001 versus 408 hours for the phenyl system. In the polyester system, the alkylcarbamate system gives 504 hours of blister free coatings versus 398 hours for the phenylcarbamate system in a detergent bath.

EXAMPLES 45-52

Unpigmented coating formulations are prepared by the general procedure described above and cured on steel panels at 177°C. for 6-PC and benzoguanamine resins (CYMEL® 1123) and at 125°C. for melamine oligomer methoxymethyl (CYMEL® 303) resins, using hydroxy functional polyesters and 20 min. cure times. In addition to detergent resistance, salt spray resistance and pencil hardness are measured. The formulations used and the results obtained are set forth in Table 6, as follows:

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Table 6. Carbamylmethylmelamine-Cured Polyesters

<u>Example</u>	<u>45</u>	<u>46</u>	<u>47</u>	<u>48</u>	<u>49</u>
<u>Composition (parts by weight)</u>					
Polyester (CYPLEX®1473-5)	60	70	75	80	85
5 6-PC	40	30	25	20	15
CYMEL® 303	--	--	--	--	--
CYMEL® 1123	--	--	--	--	--
EPON® 1001	--	--	--	--	--
TBDAS	1	1	1	1	1
10 p-TSA	--	--	--	--	--
<u>Properties</u>					
Pencil hardness	3H-4H	2H-3H	2H-3H	3H	2H-3H
Knoop hardness	23	16.4	16.6	17.5	16
Reverse Impact	60	150	150	160	160
15 T-Bond	T5	T2	T4	T3	T1
Detergent Resistance,					
Blister Code, Hrs.					
48	F9	F8	M9	M9	M8
72	M8	M8	M8	M8	M8
20 216	D7	D6	D7	D7	D6
384	D7	D4	D6	D6	D4
Salt Spray Exposure					
Blister Code, 46 Hrs.					
Tape Pull, mm	0	1	1	1	0
25 168 hrs	10	10	10	10	10
mm	1	2	2	2	3
300 hrs	10	10	10	10	10
mm	2.5	2	2.5	2	4
468 hrs	10	10	10	10	10
30 mm	4	3.5	4	3.5	5

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Table 6. (Continued)

<u>Example</u>	<u>50</u>	<u>51</u>	<u>52</u>	<u>50A</u>	<u>51A</u>
<u>Composition (parts by weight)</u>					
Polyester (CYPLEX® 1473-5)	75	70	65	75	75
5 6-PC	15	20	25	--	--
CYMEL® 303	--	--	--	15	--
CYMEL® 1123	--	--	--	--	15
EPON® 1001	10	10	10	10	10
TBDAS	1	1	1	--	--
10 p-TSA	--	--	--	.4	.4
<u>Properties</u>					
Pencil hardness	2H-3H	3H-4H	4H-5H	2H-3H	H-2H
Knoop hardness	16	21.5	19.4	15.6	15.5
Reverse Impact	160	160	160	160	150
15 T-Bond	T2	T4	T4	T2	T4
Detergent Resistance, Blister Code, Hrs.					
48	10	10	10	10	10
72	10	10	10	10	10
20 216	10	F9	10	(168-D9)	D7
384	10	D9	10	--	D4
Salt Spray Exposure					
Blister Code, 46 Hrs.	10	10	10	10	10
Tape Pull, mm	0	0	0	0	0
25 168 hrs	10	10	10	--	10
mm	0	0	0	--	0
300 hrs	10	10	10	--	10
mm	0	0	0	--	0
468 hrs	10	10	10	--	--
30 mm	0	0	0	--	--
1080 hrs	10	10	10	D4	F2
mm	0	0	0	0	4
1440 hrs	10	10	10	M4/6	--
mm	0	0	0	Striped	--

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EXAMPLES 53-60

Unpigmented coating formulations are prepared by the general procedure described above and cured on steel panels at 177°C. for 6-PC and at 125°C. for the 5 melamine oligomer resin, using a different hydroxy-functional polyester and a 20 min. cure time. The formulations used and the results obtained are set forth in Table 7, as follows:

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Table 7. Carbamylmethylmelamine-Cured Polyesters

	<u>Example</u>	<u>53</u>	<u>54</u>	<u>55</u>	<u>56</u>	<u>57</u>
	<u>Composition (parts by weight)</u>					
	Polyester (CYPLEX® 1473-5)	60	67	70	75	80
5	6-PC	40	33	30	25	20
	CYMER® 303	--	--	--	--	--
	EPON® 1001	--	--	--	--	--
	TBDAS	1	1	1	1	1
	p-TSA	--	--	--	--	--
10	<u>Properties</u>					
	Pencil hardness	4H	4H	4H	2H-3H	2H-3H
	Knoop hardness	16.6	16.4	16.4	15.2	14.2
	Reverse Impact	40	110	120	160	160
	T-Bond	T4	T3	T3	T3	T2
15	<u>Detergent Resistance,</u>					
	Blister Code, Hrs.					
	48	F9	F9	F9	M9	F9
	72	F9	F8	F9	M8	F9
	216	D6	D6	D7	D8	D8
20	384	M/D7	D5	D7	D7	D7
	<u>Salt Spray Exposure</u>					
	Blister Code, 46 Hrs.	10	10	10	10	10
	Tape Pull, mm	0	1	0	2	1
	168 hrs	10	10	10	10	10
25	mm	0	2	2	3	2
	300 hrs	10	10	10	10	10
	mm	0	2	4	4	4
	468 hrs	10	10	10	10	10
	mm	1	4	5	6	6
30	1080 hrs	--	--	--	--	--
	mm	--	--	--	--	--
	1440 hrs	--	--	--	--	--
	mm	--	--	--	--	--

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Table 7. (Continued)

	<u>Example</u>	<u>58</u>	<u>59</u>	<u>60</u>	<u>58A</u>	<u>58B</u>
<u>Composition (parts by weight)</u>						
	Polyester (CYPELEX®1473-5)	75	70	65	75	75
5	6-PC	15	20	25	--	--
	CYMEL® 303	--	--	--	25	15
	EPON® 1001	10	10	10	--	10
	TBDAS	1	1	1	--	--
	p-TSA	--	--	--	0.4	0.4
10	<u>Properties</u>					
	Pencil hardness	3H-4H	4H	3H-4H	H-2H	2H-3H
	Knoop hardness	15.4	16.6	15.5	12.6	14
	Reverse Impact	160	160	160	150	130
	T-Bond	T2	T3	T4	T3	T5
15	Detergent Resistance, Blister Code, Hrs.					
	48	10	10	10	10	F8
	72	10	10	10	F9	F8
	216	M8	M9	F9	D7	D6
20	384	M8	M/D9	D9	D4	--
<u>Salt Spray Exposure</u>						
	Blister Code, 46 Hrs.	10	10	10	10	D8
	Tape Pull, mm	0	0	0	0	4
	168 hrs	10	10	10	D9	--
25	mm	0	0	0	2	5
	300 hrs	10	10	10	--	--
	mm	0	0	0	3	8
	468 hrs	10	10	10	--	--
	mm	0	0	0	4	10
30	1080 hrs	10	10	10	M1	--
	mm	0	0	0	disint.	disint.
	1440 hrs	10	M /	10	--	--
	mm	1	0	0	--	--

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EXAMPLES 61-68

Titanium dioxide pigmented coating formulation
are prepared by the general procedure described above
and cured on steel panels, using 6-PC and methylolmel-
5 amine and methylolbenzoguanamine resins, the latter two
as controls. The formulations used and the results
obtained are set forth in Table 8 as follows:

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Table 8. TiO₂ - Pigmented Cured Coatings

<u>Example</u>	<u>61A</u>	<u>61</u>	<u>62</u>	<u>62A</u>	<u>63A</u>	<u>63</u>
<u>Composition (parts by weight)</u>						
JONCRYL®500 ^a	--	--	75	75	--	--
5 AROPLAZ 1710 R60 ^b	75	75	--	--	--	--
CYMEL® 1473-5 ^c	--	--	--	--	--	--
CARGILL 5775 ^d	--	--	--	--	75	75
CYMEL® 303 ^c	25	--	--	--	15	--
CYMEL® 1123 ^c	--	--	--	25	--	--
10 CYMEL 6-PC ^e	--	25	25	--	--	25
EPON® 1001 ^f	--	--	--	--	10	--
T-12 ^g	--	--	--	--	--	--
TBDAS ^h	--	1	1	--	--	1
p-TSA ⁱ	0.4	--	--	0.5	0.4	--
15 Cure Temp °C.	125	177	177	177	125	177
<u>Properties</u>						
MEK Resist	200+	200+	200+	200+	200	200+
Cross-Hatch Adhesion	3	5	4	5	5	5
Knoop hardness	16.6	17.5	26	18	10.8	21
20 Reverse Impact	5	30-40	5	5	160	40
Film Thickness, mils	1.2	1.1	1.1	1.3	1.3	1.2

a S. C. Johnson & Son, Inc., hydroxyfunctional polyacrylate;

b Spencer Kellog, Div. of Textron, Inc., siliconized
polyester;

25 c American Cyanamid Company;

d Cargill Co.

e Hexa(propylcarbamylmethyl)melamine (Ex. 1);

f Shell Chemical Co.;

g Dibutyltin dilaurate;

30. h Tetrabutyl diacetoxystannoxane;

i p-toluenesulfonic acid;

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Table 8. (Continued)

<u>Example</u>		<u>61A</u>	<u>61</u>	<u>62</u>	<u>62A</u>	<u>63A</u>	<u>63</u>
	Detergent Resistance, Blister Code, Hrs.						
5	48	10	10	10	D8	10	D9
	96	M/D8/9	D8	10	D6	M9	D9
	124	--	--	10	D9	M	
	148	--	--	10	VD6	VD9	VD9
	192	D8	D6	10	D4	peeled	VD9
10	288	VD6	D4	D9	D2		VD4
	336	--	--	D9	D2		
	Salt Spray Exposure						
	Blister Code, 46 Hrs.	10	10	10	D9	10	10
	Tape Pull, mm	1	0	0	0	0	0.5
15	336 hrs	10	D8	10	D9	10	10
	mm	2	1.5	0	3	1	0
	800 hrs	10	D8	10	D9	10	10
	mm	5	5	0	11	8	0

NC = no change

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The foregoing examples show that even without the epoxy resin (EPON® 1001) additive, the alkylcarbamate melamine systems outperform all state of the art melamine systems, which include the phenyl carbamate, alkoxyethyl and benzoguanamine systems in detergent resistance. The above-mentioned results are unexpected because alkyl urethanes on heating are known to be poorer leaving groups than the phenoxy blocked isocyanate system and therefore they would be expected to give somewhat inferior coating properties. The foregoing examples also show that the alkylcarbamylmethylated melamines of this invention also provide coatings with outstanding salt spray resistance in comparison with other melamine based systems.

It is generally the case also, that the alkylcarbamylmethylated melamines of this invention afford coatings with good color stability. While the phenyl analog gives off white to tan coatings on cure, the alkyl systems are unchanged. In addition, the phenylcarbamate based coatings on exposure to U.V. light change to a much darker color (tan to light brown), while the alkylcarbamate systems change only slightly to a light tan color. Finally, the data show that while outstanding resistance properties and color stability have been obtained with the alkylcarbamate melamines, other important and desirable coatings properties such as Knoop hardness, impact and solvent resistance (MEK rubs + 200) have been maintained as is the case with conventional resins.

EXAMPLE 69

The product of Example 1 can be used as a binder for foundry sand. The binder as an acetone solution (e.g., 77 wt. %) containing 1 part/100 of TBDSA catalyst is kneaded with sand and the solvent is vaporized by heating. The amount of binder to sand is preferably 1-5.5 parts to

35 core mold and heated at 200-300°C. for 30 sec. to two

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minutes. A core having good strength and showing little tendency to give off gases having a strong smell (e.g., formaldehyde) will be obtained.

Instead of sand, glass powder and glass fiber can 5 be substituted, in which case, thermally insulating shapes having good structural integrity will be obtained.

The above-mentioned patents and publications are incorporated herein by reference. Many variations of this invention will suggest themselves to those skilled in this 10 art in light of the above, detailed description. Instead of using n-propylcarbamylmethylated- and methylcarbamylmethylated melamines as curing agents in the formulations of Tables 1-8, the corresponding alkyl (and mixed alkyl) carbamylmethylated melamine and melamine oligomers of 15 Examples 3-16 can be used. Instead of tetrabutyldiacetoxy stannoxtane and, dibutyltin dilaurate as cure catalysts, lead octoate, and stannous octoate can be used. Instead of hydroxyfunctional polyesters and polyacrylates, epoxy resins, such as the polyglycidylethers of bisphenol A and 20 the reaction products thereof with amines and ammonia can be used. All such obvious modifications are within the full intended scope of the appended claims.

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CLAIMS:

1. A triazine compound selected from
 - (i) a triaminotriazine compound of the formula $C_3N_6(CH_2OR)^{6-x}(CH_2NHCOOR^1)_x$;
 - (ii) a benzoguanamine compound of the formula $C_3N_5(C_6H_5)(CH_2OR)^{4-y}(CH_2NHCOOR^1)_y$;
 - (iii) an oligomer of (i) or of (ii); and
 - (iv) a mixture of at least two of any of (i), (ii) and (iii), wherein the R groups are, independently, hydrogen or alkyl from 1 to 12 carbon atoms, the R¹ groups are, independently, alkyl of from 1 to 20 carbon atoms, x is in the range of from about 2 to about 6, and y is in the range of from about 2 to about 4.
2. A triaminotriazine compound (i) as defined in Claim 1 wherein x is from about 2.8 to about 6, and wherein R is C₁-C₈ alkyl and R¹ is methyl, ethyl, n-propyl, butyl, i-propyl, n-octyl, 2-ethylhexyl, n-octadecyl, or a mixture of any of the foregoing.
3. A benzoguanamine compound (ii) as defined in Claim 1 wherein x is from about 2.2 to about 4, and wherein R is C₁-C₈ alkyl and R¹ is methyl, ethyl, n-propyl, butyl, i-propyl, n-octyl, 2-ethylhexyl, n-octadecyl, or a mixture of any of the foregoing.
4. An oligomer of a triaminotriazine compound (iii) (i) as defined in Claim 1 wherein R is C₁-C₈ lower alkyl and R¹ is methyl, ethyl, n-propyl, butyl, i-propyl, n-octyl, 2-ethylhexyl, n-octadecyl, or a mixture of any of the foregoing.
5. A curable composition comprising:
 - (a) an active hydrogen-containing material;
 - (b) a triazine compound selected from
 - (i) a triaminotriazine compound of the formula $C_3N_6(CH_2OR)^{6-x}(CH_2NHCOOR^1)_x$;
 - (ii) a benzoguanamine compound of the formula $C_3N_5(C_6H_5)(CH_2OR)^{4-y}(CH_2NHCOOR^1)_y$.

(iii) an oligomer of (i) or of (ii); and
(iv) a mixture of at least two of any of (i),
(ii) and (iii), wherein the R groups are, independently,
hydrogen or alkyl from 1 to 12 carbon atoms, the R¹ groups are,
independently, alkyl of from 1 to 20 carbon atoms, x is
in the range of from about 2 to about 6, and y is in the range
of from about 2 to about 4; and

(c) a cure catalyst.

6. A curable composition as defined in Claim 5
wherein the active hydrogen-containing material is a
polymeric material containing at least one class of reactive
groups selected from carboxyl groups, alcoholic hydroxy
groups, amide groups, amine groups or a mixture of any of
such groups or a group convertible to any of such groups.

7. A curable composition as defined in Claim 5
wherein the triazine compound is a triaminotriazine compound
(i) wherein x is from about 2.8 to about 6, R is C₁-C₈ alkyl
and R¹ is methyl, ethyl, n-propyl, butyl, i-propyl,
n-octyl, 2-ethylhexyl, n-octadecyl, or a mixture of any
of the foregoing.

8. A curable composition as defined in Claim 5
wherein the cure catalyst comprises tetrabutyldiacetoxy
stannoxyane, dibutyltin dilaurate or dimethyltin dilaurate.

9. A substrate protectively coated with a cured
composition as defined in Claim 5.

10. An article of manufacture comprising a cured
composition as defined in Claim 5 and a filler.

